

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
7 December 2000 (07.12.2000)

PCT

(10) International Publication Number  
WO 00/72767 A1

(51) International Patent Classification<sup>7</sup>: A61B 17/58

(21) International Application Number: PCT/US00/14840

(22) International Filing Date: 26 May 2000 (26.05.2000)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
09/321,369 27 May 1999 (27.05.1999) US

(71) Applicant and

(72) Inventor: PHILLIPS, Jonathan [US/US]; 3527 Bellington Drive, Orlando, FL 32835 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

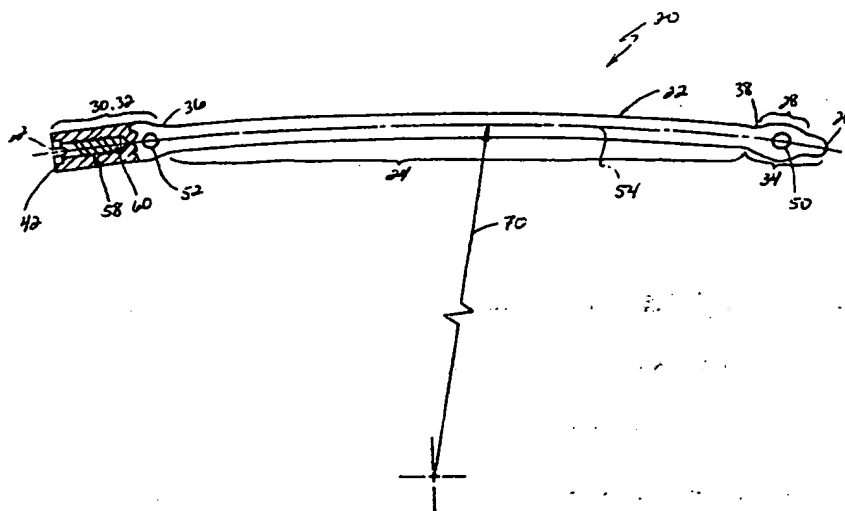
Published:

— With international search report.

(74) Agents: ROONEY, Kevin, G. et al.; Wood, Herron & Evans, L.L.P., 2700 Carew Tower, Cincinnati, OH 45202 (US).

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: PEDIATRIC INTRAMEDULLARY NAIL AND METHOD



(57) Abstract: An intramedullary nail (20) and related method for fixing a fracture in a long bone. The nail (20) comprises an elongate member (22) having a longitudinal axis (54), a proximal end section (32), a distal end section (34) and a solid central section (24) extending between the proximal and distal end sections (32, 34). The proximal and distal end sections (32, 34) respectively include proximal and distal fastener receiving areas (28, 30) of greater cross section dimensions than the central section (24). Each fastener receiving area (28, 30) includes at least one hole (50, 52) extending transverse to the longitudinal axis for receiving a cross fastener (110) adapted to secure to the bone on opposite sides of the elongate member (22). The proximal and distal end sections (32, 34) thereby provide rigid anchoring locations relative to the central section (24) and the central section provides flexibility to promote healing of the fracture.

WO 00/72767 A1

THIS PAGE BLANK (USPTO)

## **PEDIATRIC INTRAMEDULLARY NAIL AND METHOD**

This application is a continuation-in-part of U.S. Application Serial No. 09/321,369, filed May 27, 1999, now pending, the disclosure of which is entirely incorporated by reference herein and the benefit of which is claimed under 35 U.S.C. § 120.

### **BACKGROUND OF THE INVENTION**

#### **Field of the Invention**

The present invention relates to devices and methods for repairing bone fractures, and, more particularly, to intramedullary nails and related internal fixation methods especially suitable for repairing long-bone fractures in children.

#### **Description of Related Art**

The use of intramedullary nails in the repair of long-bone fractures, such as in the femur, has been known in the orthopedic field. Exemplary devices include those known as the Rush and Enders and the Kuntschner nails, as well as those disclosed in U.S. Pat. Nos. 5,713,902 to Friedl; 5,697,930 to Itoman et al.; 5,573,536 to Grosse et al.; 5,562,666 to Brumfield; 5,374,235 to Ahrens; 5,312,406 and 5,167,663 to Brumfield; 5,122,141 to Simpson; 5,066,296, 5,041,114, and 4,776,330 to Chapman et al.; 4,976,258 to Richter et al.; 4,875,475 to Comte et al.; 4,846,162 to Moehring; 4,506,662 to Anapliotis; and 4,475,545 to Enders.

Referring to Fig. 1, a special problem in pediatric orthopedics exists in that reaming through the typical entry point in a femur 10, i.e., the piriformis fossa 11, can be too dangerous for the child. This is due to the presence of an artery 12 that supplies blood to the proximal femur.

5 Specifically, this is the lateral epiphyseal artery 12 which is a branch of the femoral artery. If this artery 12 is damaged during the fixation procedure, such as while the intramedullary canal is being reamed to accept a nail, or possibly during insertion or after insertion of the nail, various complications can result. The lateral epiphyseal artery 12 supplies 75% of the blood to  
10 the growing femoral head 16. If this artery 12 is damaged, then much of the femoral head 16 will die or necrose. The femoral head 16 will then heal with an irregular shape which inevitably leads to hip arthritis.

Various nails, such as flexible Rush nails, are non-interlocked meaning that cross fasteners are not used to secure the nail to the bone. These nails are  
15 often small diameter rods, on the order of approximately 3-4 mm in diameter. In addition to being flexible to a significant degree prior to plastic deformation, non-interlocked solid nails or rods can be relatively easily bent with plastic deformation to a desired shape. A plurality of these nails or rods are typically driven into the intramedullary canal depending on the  
20 support necessitated by the fracture and bone characteristics of the patient.

Other more rigid solid or hollow nails are interlocked to the bone using cross fasteners typically at the proximal and distal ends of the nail. Unlike non-interlocked nails, interlocked nails require sufficient cross-sectional dimensions to accommodate holes necessary for the cross fasteners.

Currently available interlocked nails can be inserted away from the lateral epiphys al artery 12 but are so rigid that they migrate during insertion dangerously close to the artery 12 and can endanger it. In addition, the large proximal size of small adult interlocked nails, which have typically  
5 been used in children, increases the potential for damage to the growth plate 17 at the proximal femur.

Among possible solutions, retrograde nailing avoids the proximal femur but also has at least one potential problem. The nails must be introduced close to the distal femoral growth plate or physis 19 (Fig. 5)  
10 at an awkward angle, potentially causing growth arrest distally on the femur, i.e., adjacent the knee. An approach through the greater trochant r 18 is also well recognized, but usually only one small diameter non-interlocked nail or rod can be used because of the narrow safe entry zone of the greater trochanter. A second small diameter nail or rod needs to be  
15 inserted retrograde or through the opposite end of the femur in these situations. These small diameter, flexible nails allow flexure after insertion and the slightly added stress to the bone allowed by this flexure promotes faster bone healing. These non-interlocked nails work well for transverse fractures. However, spiral or comminuted fractures often need additional  
20 external support, such as with a cast or brace. This is due to the inability of the non-interlocked nail to effectively prevent rotation or length compromise at the fracture.

It would therefor be desirable t pr vide an int rlocked intramedullary nail, especially suitabl for pediatric use, which provides

flexibility along a majority of the length of the nail to facilitate faster healing of a fracture, but which also provides for secure interlocking of the nail to the bone with cross fasteners to prevent compromising the fracture due to rotation or shortening at the fracture site. Ideally, such a nail and related methods of insertion would minimize trauma to the growth plates of the femur as well as the arteries that supply blood to the proximal end of the femur while still allowing easy insertion and fixation within the intramedullary canal.

### **SUMMARY OF THE INVENTION**

10 In one general aspect, the present invention provides an intramedullary nail for insertion within an intramedullary canal of a long bone and fixing a fracture in the long bone. The nail is especially suitable for adolescent or preadolescent aged children, however, the nail may be useful in other orthopedic applications as well. The nail generally comprises

15 an elongate member having a longitudinal axis, a proximal end section, a distal end section and a solid central section extending between the proximal and distal end sections. The proximal and distal end sections respectively have fastener receiving areas of greater cross sectional dimensions than the central section. The fastener receiving areas each

20 include at least one hole extending transverse to the longitudinal axis of the elongate member for receiving a cross fastener adapted to secure to the bone on opposite sides of the elongate member. The proximal and distal end sections provide rigid anchoring locations relative to the central section

and the central section provides flexibility to promote healing of the fracture.

In the preferred embodiment, the central section of the elongate member is curved in the sagittal plane to generally follow the curvature of a femur. The proximal and distal end sections are bent out of this plane and form acute angles with respect to the sagittal plane. The proximal and distal end sections are each bent laterally to one side of the central section. The side to which the proximal and distal end sections are bent depends on whether the nail will be used in a right or left femur and also allows easier insertions across the fracture. The bend of the distal end section allows for easier insertion of the nail from an insertion point extending through the greater trochanter of the femur. The bend of the proximal end section ensures that the proximal tip is presented directly at the insertion point after fixation so that it may be easily accessed for removal purposes upon healing of the bone.

In general, a method of fixing a fracture in a long bone of a patient, in accordance with the invention, includes providing an elongate member having a solid central section with a cross sectional dimension and having proximal and distal fastener receiving areas of increased cross sectional dimension relative to the cross sectional dimension of the central section. The fastener receiving areas each have at least one hole extending transverse to a longitudinal axis of the elongate member. The method involves inserting the elongate member into the intramedullary canal of the long bone through an insertion point and across the fracture and inserting

cross fasteners through each of the holes and into the bone on opposite sides of the elongate member to fix the fracture of the long bone against rotational and lengthwise movements.

The bone nail and method of this invention allow the surgeon

5 to better avoid the critical arterial blood supply to the femoral head which crosses at the piriformis fossa, i.e., the traditional point of entry for an adult nail. Instead, the entry point is on the greater trochanter, a location distinctly lateral of the critical region of the piriformis fossa. The very vulnerable growth plate between the greater trochanter and the femoral

10 neck is also avoided using the nail and entry point in accordance with this invention. Another unique feature of the invention is the ability of the bone nail to be flexible, yet custom bent to match the exact geometry of the proximal femur, while also allowing interlocking with cross fasteners. In general, the preferred inventive nail provides interlockability, a relatively

15 small solid cross section to provide flexure along at least the majority of the length of the nail, and malleability to allow custom bending especially at the proximal end.

The features that characterize the invention, both as to organization and method of operation, together with further objects,

20 features and advantages thereof, will be better understood from the following written description taken in conjunction with the accompanying drawings. It is to be expressly understood that the drawings and detailed description thereof are for the purpose of illustration and description and is not intended as a definition of the limits of the invention. These and other



objects attained, and advantages offered, by the present invention will become more fully apparent as the description that now follows is read in conjunction with the accompanying drawings.

### **BRIEF DESCRIPTION OF THE DRAWINGS**

5                    Fig. 1 is a perspective view of a portion of a femur including the femoral head and the greater and lesser trochantic regions.

Fig. 2 is a side elevational view of an intramedullary nail of the present invention shown in the anterior-posterior plane.

10                   Fig. 3 is an elevational view of the nail shown in Fig. 2, but illustrating proximal and distal bends.

Fig. 4 is an elevational view of the nail shown in Fig. 3, but rotated 90°.

Fig. 5 is a schematic view of the step of initially inserting the bone nail of Figs. 3 and 4 into a femur.

15                   Fig. 6 is a schematic view similar to Fig. 5, but illustrating the nail further inserted across the bone fracture.

Fig. 7 illustrates a drill rig and fastener inserting rig being used to cross fasten the nail after full insertion.

20                   Fig. 8 is a schematic illustration of the fully inserted and cross fastened bone nail.

Fig. 8A is a fragmented, partially sectioned view of the distal end section of the bone nail and the cross fastener of the present invention.

Fig. 9 is an elevational view showing one illustrative bending device being used to bend the proximal end section of the bone nail.

### DETAILED DESCRIPTION

An exemplary intramedullary nail 20, as illustrated in FIGS. 2-4, comprises an elongate member 22. In an embodiment intended for use in a femur of a child or adolescent, typically ages 6-14, for example, the nail 20 is formed from titanium and has a generally cylindrical shape with a diameter along a solid central section 24 of between 4mm and 7mm, although this is not intended as a limitation. Central section 24 should have a certain amount of elastic flexibility to permit accurate placement within the femoral canal as discussed below. Such elastic flexibility, which permits some flexure during movement and weight bearing activity, confers an additional benefit in that it has been found to stimulate bone healing. The nail 20 preferably ends in a rounded, smooth and tapered distal tip 26.

15 The nail 20 has two widened sections or fastener receiving areas 28, 30 in respective proximal and distal end sections 32, 34. Each fastener receiving area 28, 30 is formed with an increased cross sectional dimension relative to central section 24. Also, smooth, gradual transitions 36, 38 with the central section 24 avoid sharp edges along the length of nail 20. The distal fastener receiving area 28 is positioned adjacent to but in spaced relation from distal tip 26. The distal fastener receiving area 28 tapers on both proximal and distal sides thereof. The proximal fastener receiving area 30 extends to proximal tip 42; that is, there is a distal taper

but no proximal taper, and the widened proximal fastener receiving area 30 instead continues to the proximal tip 42 and forms the entire proximal end section 32.

Each fastener receiving area 28, 30 includes a generally  
5 cylindrical hole 50, 52 extending generally normal to the portion of the longitudinal axis 54 in which it is located for receiving a fastener (not shown). Holes 50, 52 preferably have diameters in a range of 3 mm to 4 ½ mm.

Proximal tip 42 includes attachment structure 58 for receiving  
10 a driver, described below, which may be a conventional driver used in the bone nail art. Preferably, attachment structure 58 comprises a threaded axial bore 60 extending along axis 54 and engageable with an externally threaded driver. A notch 62 extends across bore 60 and, as is known, aligns the driver for purposes of later drilling and cross fastening nail 20, as  
15 will be discussed below. As further shown in Figs. 2 and 4, nail 20 is curved in an anterior direction along a radius of curvature 70 to generally conform nail 20 to the typical femoral curvature. Preferably, this radius of curvature 70 is in the range of 30 inches to 60 inches. A proximal bend 72 and a distal bend 74 are formed, respectively, in central section 24 directly  
20 adjacent proximal and distal end sections 32, 34. These bends 72, 74 are made in the same direction, i.e., laterally to one side of nail 20 or the other as shown best in Fig. 3. The solid lines in Fig. 3 illustrate lateral bends 72, 74 in the sagittal plane and generally in the coronal plane at angles A, B for inserting nail 20 into a left femur. Angles A', B' correspond to the

respective lateral bends of proximal and distal end sections 32, 34, shown in phantom, facilitate use of nail 20 in the right femur. As a unique feature of this invention, interlocking nail 20 may be custom bent by the surgeon just prior to use, not only to facilitate insertion in the right or left femur, but  
5 also to accommodate other particular shapes necessary for a particular patient. For example, some patients may have deformities necessitating one or more corrective osteotomies or fractures made by the surgeon. These osteotomies may also be fixed using nail 20.

In the preferred embodiment, nail 20 is formed from titanium,  
10 although other materials such as those known in the art may be used as well. Central section 24 is of solid cross section and at least substantially constant diameter with a smooth outer surface to facilitate removal in 6-9 months. Proximal and distal end sections 32, 34 are also solid, except for holes 50, 52, bore 60 and notch 62. As appreciated from Fig. 2, the axes  
15 of holes 50, 52 are coplanar. In order to provide the desired flexibility of central section 24, while retaining the cross fastening feature of the invention, fastener receiving areas 28, 30 have a cross sectional dimension greater than the cross sectional dimension of central section 24. In general, central section 24 may be formed with a solid cylindrical cross section  
20 having a diameter of between 4 mm and 7 mm. Fastener receiving area 28 has a generally bulbous, rounded shape, while proximal fastener receiving area 30, which is preferably continuous with distal end section 32, has a cylindrical cross sectional shape. At the cross section taken along the axes of respective holes 50, 52, the respective ratios of the cross sectional

dimensions at these locations is at least about 1.3:1 relative to the cross sectional dimension of central section 24. In exemplary embodiments of the invention, a nail 20 substantially as shown in Fig. 2, had a cross sectional dimension of 5.5 mm for central section 24, an 8.5 mm cross sectional dimension for proximal end section 32, and also a maximum 8.5 mm cross sectional dimension at the largest diameter portion of distal fastener receiving area 28. This nail was used on children generally weighing less than 100 lbs. For children weighing more than 100 lbs., the 8.5 mm dimensions at the proximal and distal fastener receiving areas remained the same, while the cross sectional dimension for central section 24 was increased slightly to 6.5 mm to provide additional strength but still provide the desirable flexure.

The insertion and fixation techniques according to the preferred embodiment of the invention are best illustrated in Figs. 5-8. In accordance with the invention, an insertion point 80 is created in the greater trochanter 18 of the femur 10 of a child, for example, at a distinctly lateral position relative to the piriformis fossa 11. Intramedullary canal 10a of femur 10 is drilled and reamed in a known manner to accept bone nail 20. A driver 82 is coupled with proximal tip 42, also in a known manner, and the surgeon impacts the head 84 of the driver 82 while holding the handle portion 86. Bend 74 facilitates better positioning of distal tip 26 upon insertion of nail 20 by allowing distal tip 26 to naturally follow the intramedullary canal 10a relative to the angle of the insertion point 80 in the greater trochanter 18. As shown in Fig. 6, bone nail 20 may be rotated in either direction

represented by arrow 90 as the distal tip 26 approaches and crosses the fracture 92. This allows the surgeon, using fluoroscopy, to more easily locate and enter the intramedullary canal 10a of the distal bone segment 94. After the bone nail 20 is fully inserted, as shown in Fig. 7, a drill rig 100 is attached to the proximal tip 42 through securement to handle portion 86. The drill rig 100 aligns a drill guide 102 with the proximal fastener receiving hole 52 and a drill (not shown) is used to form a hole through femur 10 in line with hole 52. A screw driving mechanism 104 is then used to insert a cross fastener 110 at this location. Using conventional fluoroscopy techniques, a second hole is drilled and a second cross fastener 112, preferably of the same design as fastener 110, is inserted through the distal hole 50, as shown in Figs. 8 and 8A. More specifically, and as represented by distal fastener 112 in Fig. 8A, fastener 112 comprises a drive head 150, a proximal threaded portion 152, a distal threaded portion 154, and a central unthreaded portion 156 which is received within hole 50. Threaded portions 152, 154 are securely engaged within cortical layer 10b of femur 10. In this manner, bone nail 20 is interlocked to femur 10 at proximal and distal locations thereby preventing undesirable rotational and/or lengthwise bone movements at the fracture site. In this interlocked or fixed position, proximal tip 42 is presented directly at the insertion point 80 on the greater trochanter 18 so that, upon healing of the fracture, the cross fasteners 110, 112 may be removed and threaded bore 60 may be engaged to withdraw nail 20 from intramedullary canal 10a.

Although bends 72, 74 may be pre-made by a manufacturer of nail 20, for example, the present invention further contemplates a manual bending device as shown in Fig. 9. Using this device, nail 20 may be placed by the surgeon in a bending device 118 jaw structure comprising three rollers 120, 122, 124 with one roller 120 acting as a fulcrum and two opposite rollers 122, 124 applying forces in the direction of arrows 130, 132. When the handles 134, 136 of the device are squeezed together in the direction of arrows 138, 140, proximal end section 32 will be bent relative to central section 24 to form bend 72 as best illustrated in Fig. 3 and as previously described. The same procedure may be used by the surgeon to bend distal end section 34 just prior to insertion within intramedullary canal 10a. This aspect of the invention allows the surgeon to custom bend these or other portions of the nail 20 to suit the needs of a particular patient prior to or during surgery.

It may be appreciated by one skilled in the art that additional embodiments may be contemplated, including similarly designed nails for fixing fractures in other long bones of child and adolescent patients. It can also be contemplated that similar nails could be useful for animal long-bone fractures.

In the foregoing description, certain terms have been used for brevity, clarity, and understanding, but no unnecessary limitations are to be implied therefrom beyond the requirements of the prior art, because such words are used for descriptive purposes herein and are intended to be broadly construed. Moreover, the embodiments of the apparatus illustrated

and described hereinafter by way of example, and the scope of the invention is not limited to the exact details of construction.

Having now described the invention, the construction, the operation and use of preferred embodiment thereof, and the advantageous  
5 new and useful results obtained thereby, the new and useful constructions, and reasonable mechanical equivalents thereof obvious to those skilled in the art, are set forth in the appended claims.



**What is claimed is:**

1. An intramedullary nail for insertion within an intramedullary canal of a long bone and fixing a fracture in the long bone, the nail comprising:
  - 5 an elongate member having a longitudinal axis, a proximal end section, a distal end section and a solid central section extending between said proximal and distal end sections, said proximal and distal end sections respectively including proximal and distal fastener receiving areas of greater cross-sectional dimensions than said central section, said fastener receiving  
10 areas each having at least one hole extending transverse to the longitudinal axis for receiving a cross fastener adapted to secure to the bone on opposite sides of said elongate member, said proximal and distal end sections thereby providing rigid anchoring locations relative to said central section, and said central section providing flexibility to promote healing of  
15 the fracture.
2. The intramedullary nail of claim 1, wherein said central section is malleable to allow bending of at least the proximal end section with respect to the central section.
3. The intramedullary nail recited in claim 1, wherein the central  
20 section of said elongate member is curved between said fastener receiving areas in a sagittal plane transverse to respective axes of the holes, said

curved central section thereby adapted to conform with the intramedullary canal.

4. The intramedullary nail of claim 3, wherein said elongate member further includes a proximal bend located distally of the fastener receiving area of said proximal end section, said proximal bend forming an acute angle relative to the sagittal plane containing the axis of curvature of said central section.

5. The intramedullary nail of claim 4, wherein said elongate member further includes a distal bend located proximally of the fastener receiving area of said distal end section, said distal bend forming an acute angle relative to the sagittal plane containing the axis of curvature of said central section and being on the same side of the sagittal plane as the proximal bend.

6. The intramedullary nail of claim 1, wherein the ratio of the cross sectional dimensions of the respective proximal and distal fastener receiving areas at the axes of said holes relative to the cross sectional dimension of said central section is at least about 1.3:1.

7. The intramedullary nail of claim 1, wherein said distal fastener receiving area is tapered on proximal and distal ends thereof and said proximal fastener receiving area is tapered on a distal end thereof.

8. The intramedullary nail of claim 1, wherein the holes extend along respective axes and the axes of said holes are generally coplanar.

9. The intramedullary nail of claim 1 further comprising cross fasteners respectively received in the holes, each cross fastener having a threaded distal tip, a threaded proximal shank and an unthreaded portion between the threaded distal tip and the threaded proximal shank, said unthreaded portion adapted to be received in one of said holes and said threaded distal tip and proximal shank adapted to engage bone matter on opposite sides of said one hole.

10. A method of fixing a fracture in a long bone of a patient having an intramedullary canal, the method comprising:

providing a elongate member having a solid central section having a cross sectional dimension and having proximal and distal fastener receiving areas of increased cross sectional dimension relative to the cross sectional dimension of the central section, the fastener receiving areas each having at least one hole extending transverse to a longitudinal axis of the elongate member,

inserting the elongate member into the intramedullary canal through an insertion point and across the fracture, and

inserting cross fasteners through each of said holes and into said bone on opposite sides of said elongate member to fix the fracture of the long bone against rotational and lengthening movements.

11. The method of claim 10, wherein at least the central section of the elongate member is curved in a sagittal plane of the patient, and further comprising:

5 prior to the inserting step, laterally bending the proximal fastener receiving area of said elongate member at an acute angle out of the sagittal plane of the patient.

12. The method of claim 11 further comprising:  
bending the distal fastener receiving area of said elongate member at an acute angle out of the sagittal plane of the patient.

10 13. The method of claim 11 further comprising:  
laterally bending the proximal fastener receiving area to conform to a right femur of the patient.

14. The method of claim 11 further comprising:  
laterally bending the proximal fastener receiving area to  
15 conform to a left femur of the patient.

15. The method of claim 11, wherein the long bone is a femur and the insertion point is a point on the greater trochanter lateral of the piriformis fossa, and the method further comprises:

laterally bending the proximal fastener receiving area to  
20 conform to the proximal femur of the patient and to present the proximal tip

of the elongate member at the insertion point for access and removal after healing of the fracture.

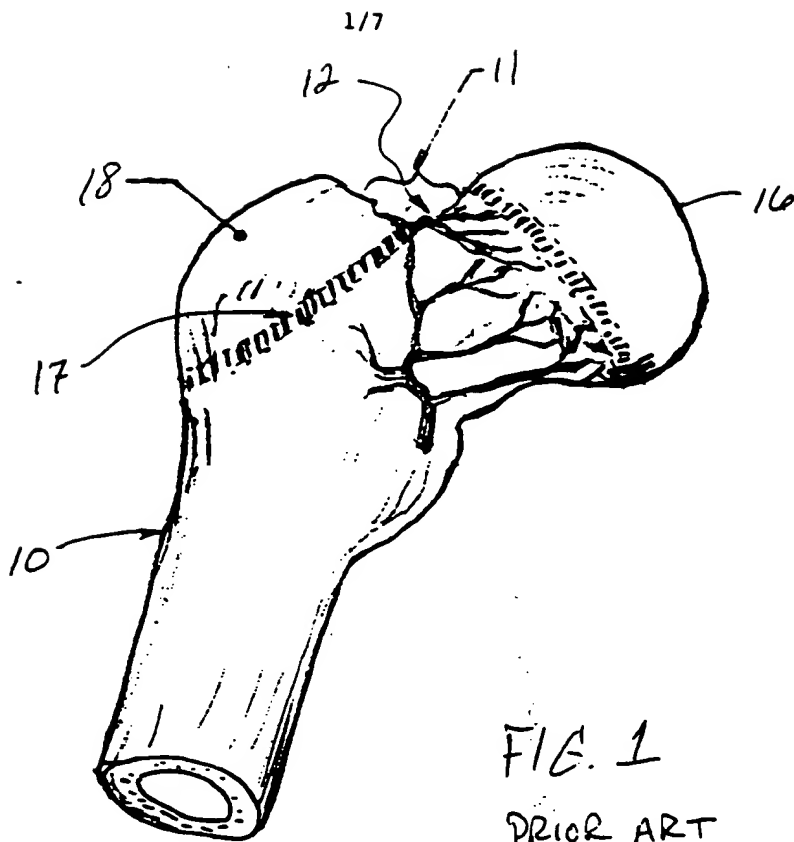
16. An intramedullary nailing system for fixing a fracture in a long bone of a patient having an intramedullary canal, the system comprising:

5 an elongate member having a longitudinal axis, a proximal end section, a distal end section and a solid central section extending between said proximal and distal end sections, said proximal and distal end sections respectively including proximal and distal fastener receiving areas of greater cross-sectional dimensions than said central section, said fastener receiving  
10 areas each having at least one hole extending transverse to the longitudinal axis for receiving a cross fastener adapted to secure to the bone on opposite sides of said elongate member, said proximal and distal end sections thereby providing rigid anchoring locations relative to said central section, and said central section providing elastic flexibility to promote  
15 healing of the fracture, and

a bending device having jaw structure configured to hold the elongate member and bend at least one of the proximal and distal end sections at angle relative to said central section.

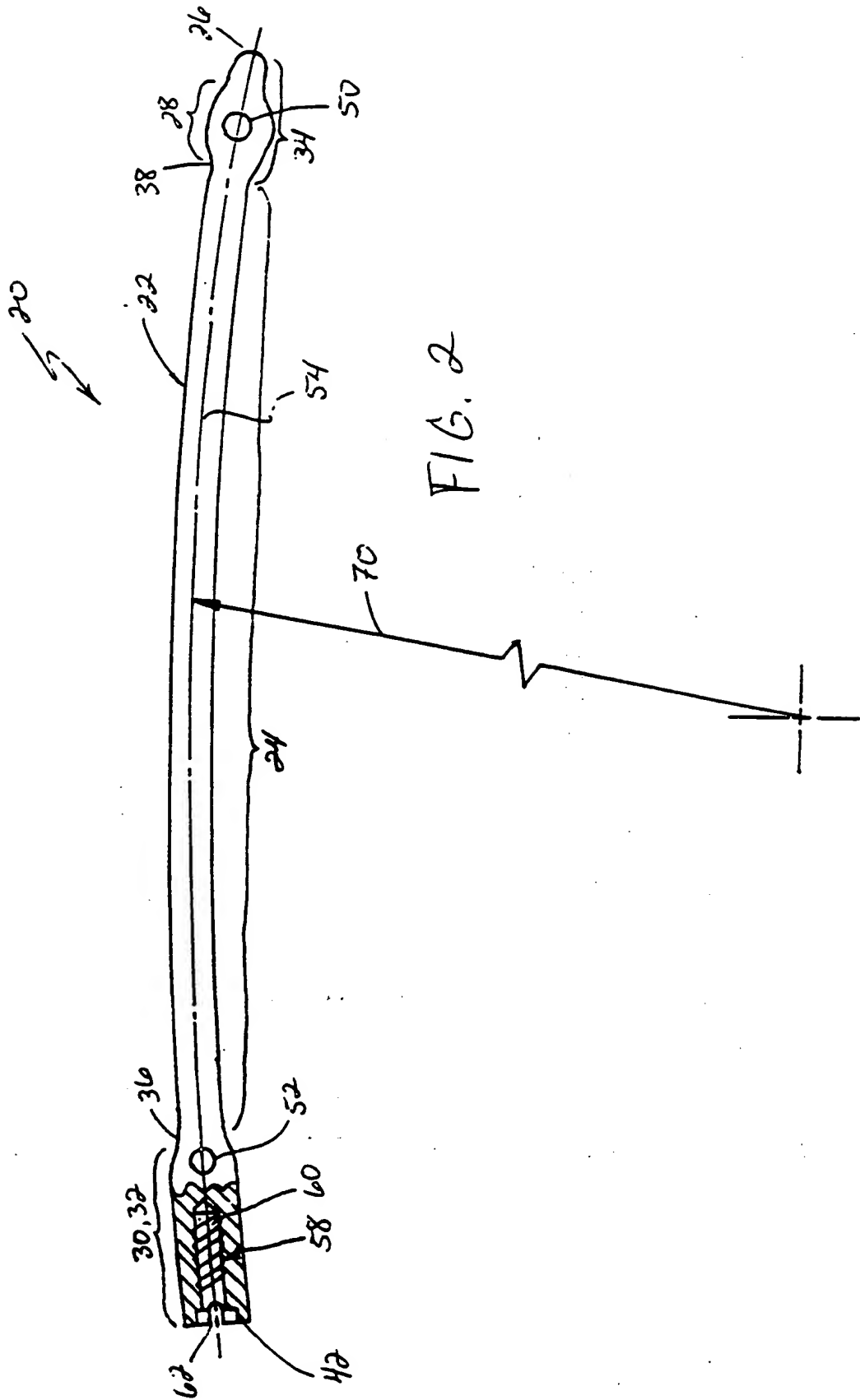
17. The system of claim 16, wherein said bending device further  
20 comprises a pair of manually operable handles coupled with said jaw structure and adapted to be squeezed together to move the jaw structure.

**THIS PAGE BLANK (USPTO)**

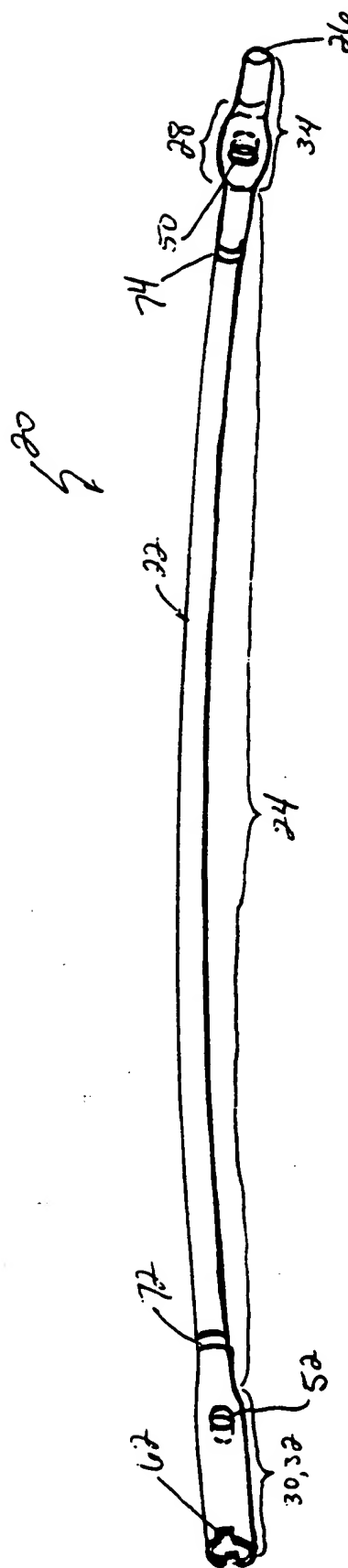
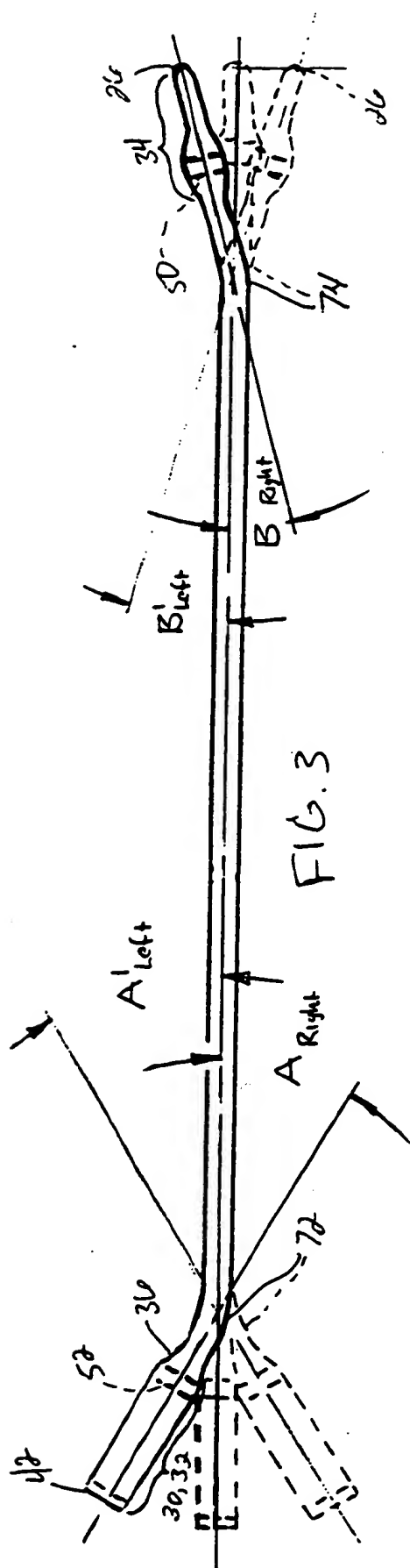


**THIS PAGE BLANK (USPTO)**

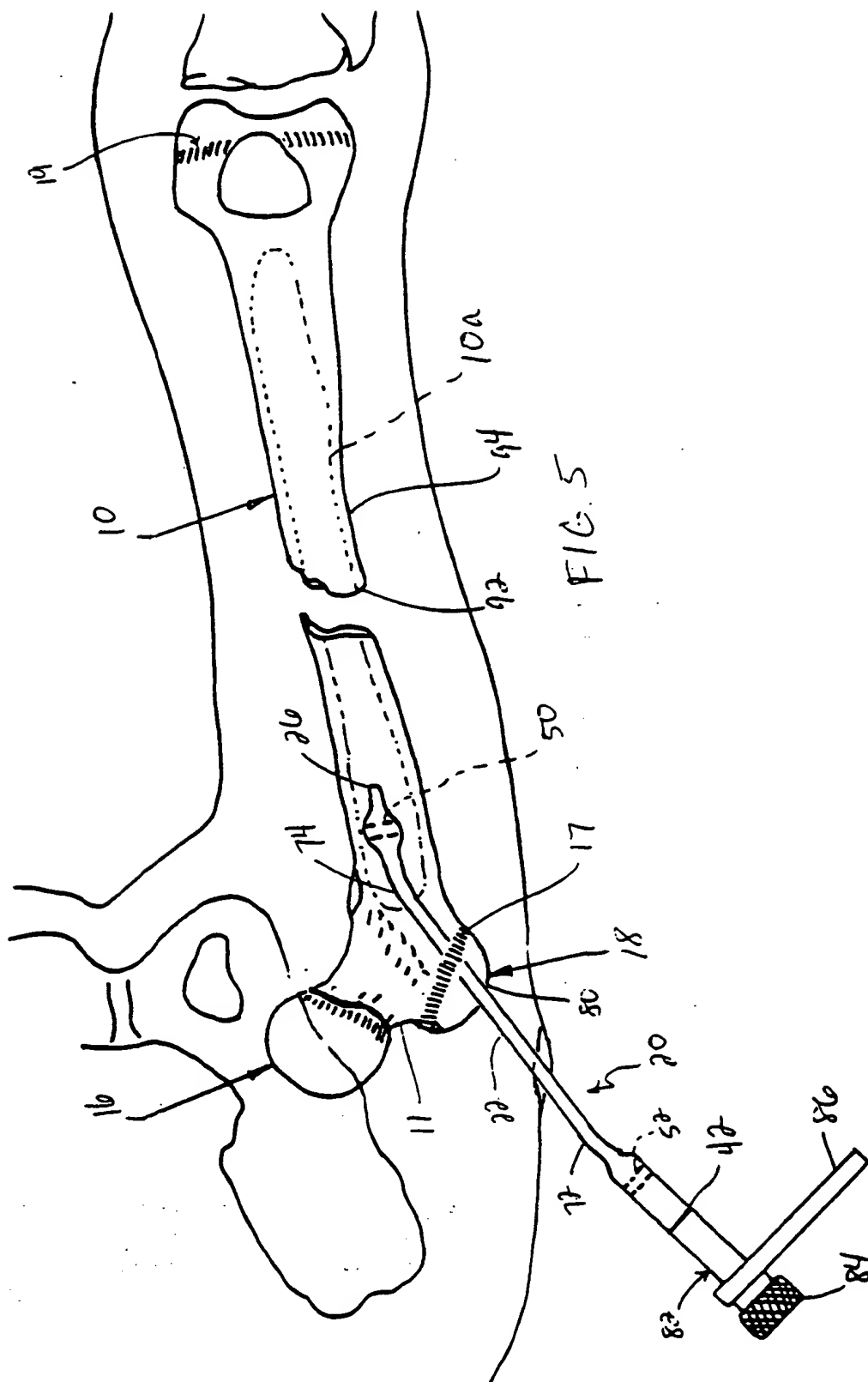




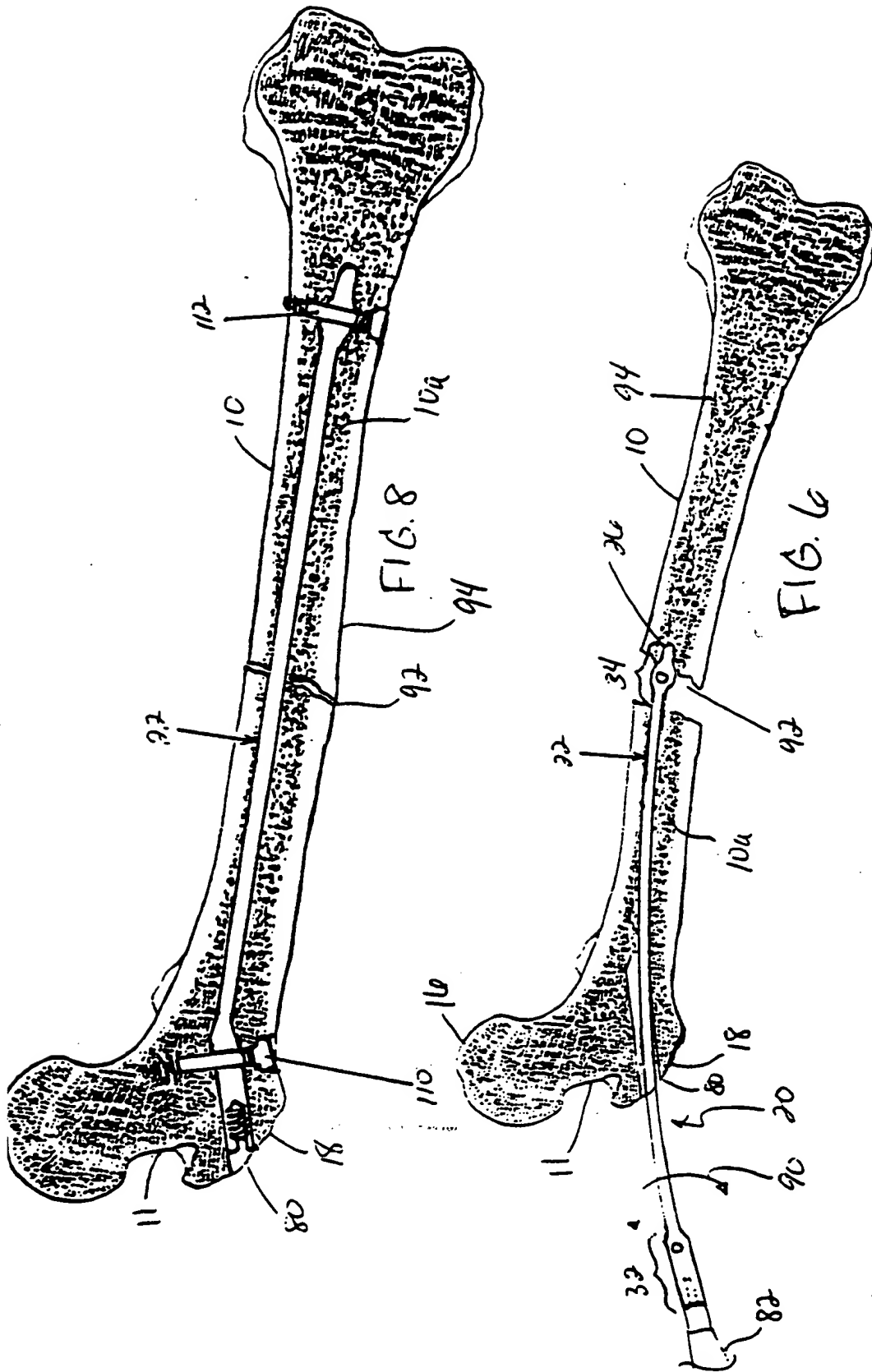
**THIS PAGE BLANK (USPTO)**



**THIS PAGE BLANK (USPTO)**

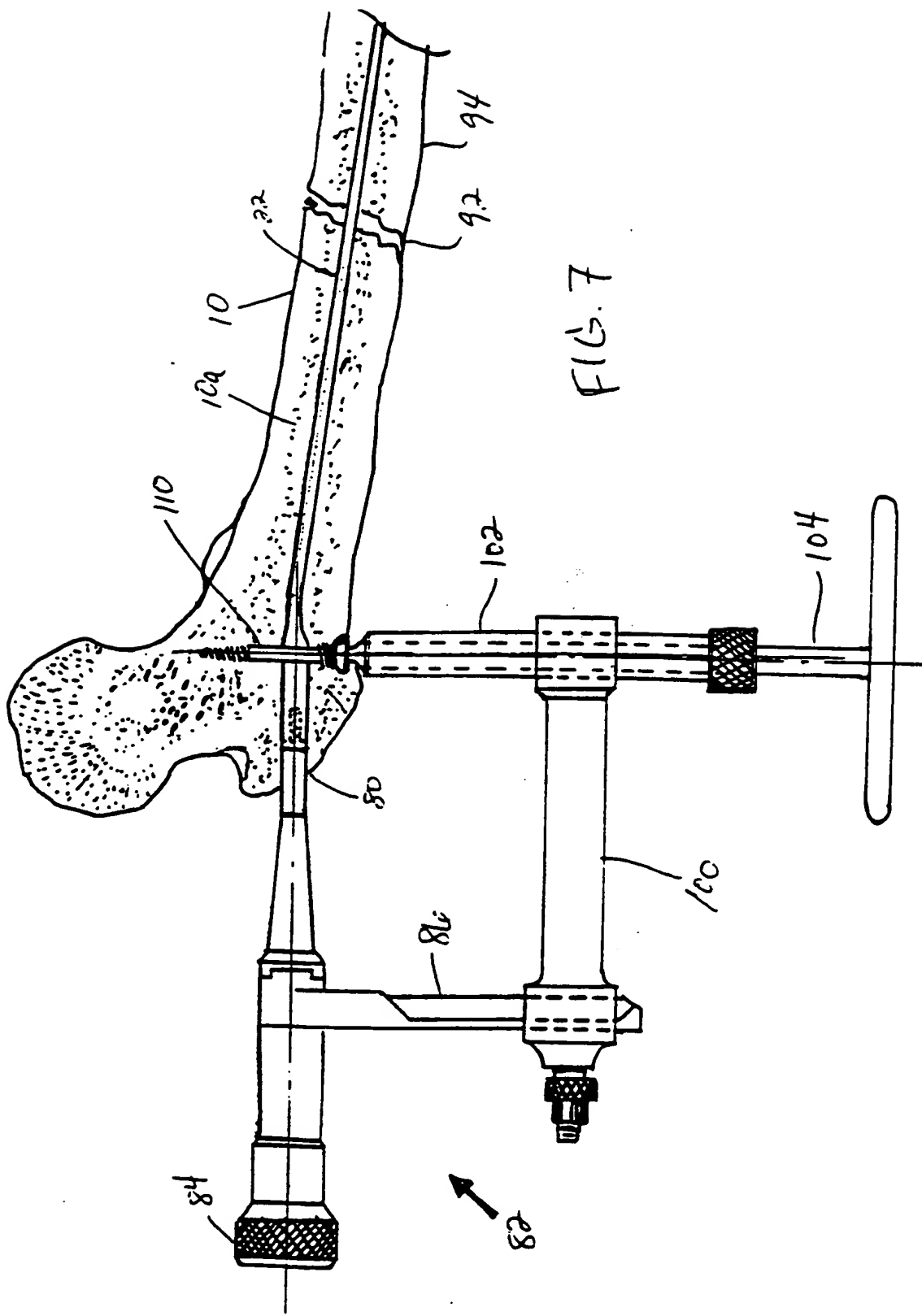


**THIS PAGE BLANK (USPTO)**

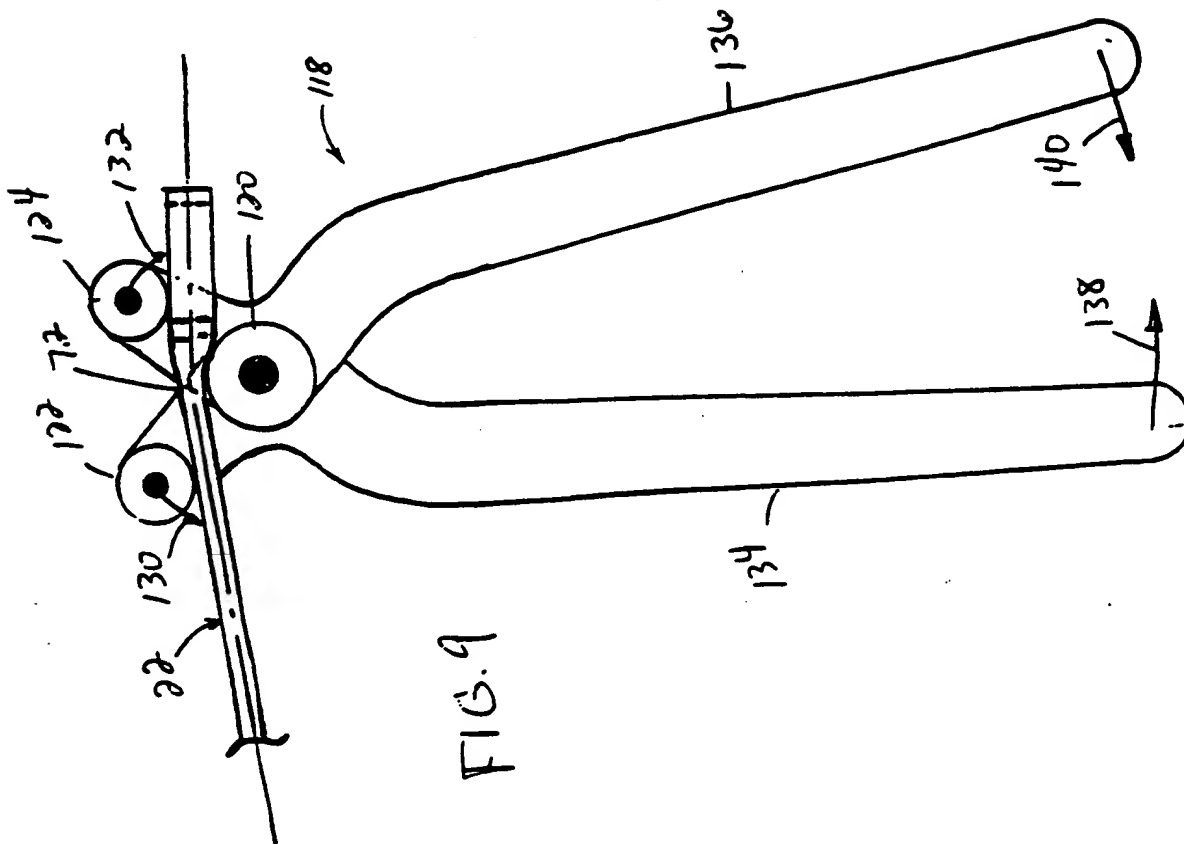
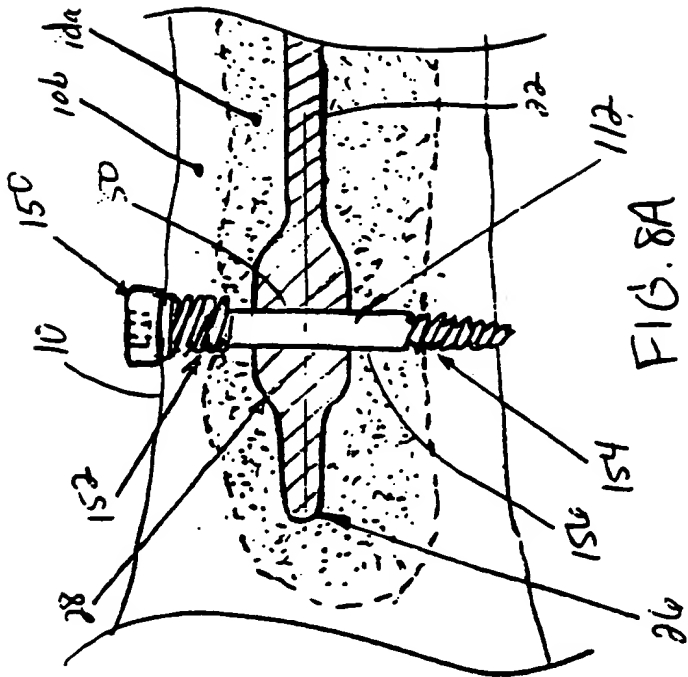


**THIS PAGE BLANK (uspto)**





**THIS PAGE BLANK (USPTO)**



**THIS PAGE BLANK (USPTO)**

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US00/14840

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61B 17/58; A61F 5/46

US CL : 606/62, 101

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 411/383, 389; 606/62-68, 72, 73, 75-77, 95, 101, 104

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
WEST

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,034,013 A (KYLE et al.) 23 July 1991, entire document.	1-8, 10-15
Y		9, 16, 17
Y	US 5,019,079 A (ROSS) 28 May 1991, entire document.	9
Y	US 4,862,883 A (FREELAND) 05 September 1989, entire document.	16, 17
A	US 5,041,115 A (FRIGG et al.) 20 August 1991, entire document.	1-17
A	US 5,035,697 A (FRIGG) 30 July 1991, entire document.	1-17

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents	* T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A* Document defining the general state of the art which is not considered to be of particular relevance	* X* document of particular relevance, the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* E* earlier document published on or after the international filing date	* Y* document of particular relevance, the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
* L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	* &* document member of the same patent family
* U* document referring to an oral disclosure, use, exhibition or other means	
* P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

10 JULY 2000

Date of mailing of the international search report

25 JUL 2000

Name and mailing address of the ISA/US  
Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

PEDRO PHILOGENE

Telephone No. (703) 308-2252

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US00/14840

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4,913,137 A (AZER et al.) 03 April 1990, entire document.	1-17